

13



Europäisches Patentamt
European Patent Office
Office européen des brevets

#4

11 Publication number:

0 100 195
A2

12

EUROPEAN PATENT APPLICATION

21 Application number: 83304170.0

22 Date of filing: 19.07.83

51 Int. Cl.: C 11 D 17/04, C 11 D 3/37,
C 09 G 1/16

50 Priority: 26.07.82 US 401805

43 Date of publication of application: 08.02.84
Bulletin 84/8

84 Designated Contracting States: BE DE FR GB IT NL

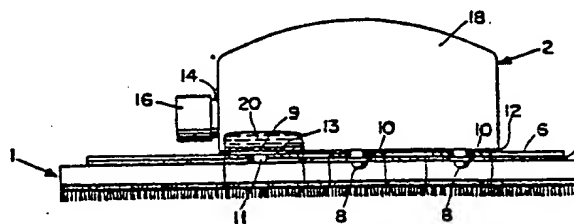
71 Applicant: THE PROCTER & GAMBLE COMPANY,
301 East Sixth Street, Cincinnati Ohio 45201 (US)

72 Inventor: Trinh, Toan, 8571 Creekwood Lane, Maineville
Ohio 45039 (US)

74 Representative: Gibson, Tony Nicholas et al, Procter
Gamble (NTC) Limited Whitley Road, Longbenton
Newcastle upon Tyne NE12 9TS (GB)

54 Car cleaner article.

57 A car cleaning article comprising a dispenser, a bristled fibrous applicator, and a substantially nonabrasive liquid cleaner composition. The article is used to clean car surfaces without an external source of water to wash or rinse.



EP 0 100 195 A2

CAR CLEANER ARTICLE

Toan Trinh

BACKGROUND OF THE INVENTIONField of the Invention

The present invention is a car cleaner article which requires no external source of water to wash or
5 rinse.

Description of the Prior Art

Car care products are numerous. Most car cleaners require large amounts of wash and rinse water. Those which do not require an external source of wash
10 and rinse water contain a hard abrasive. A number of prior art auto cleaners are disclosed in Household and Automotive Chemicals Specialties, Recent Formulations, by E. W. Flick, Noyes Data Corporation, Park Ridge, New Jersey, 1979, pp. 293-326.

15 Current car cleaners/polishes utilizing mineral-based abrasives have problems associated with their use. Such abrasives are inherently comprised of relatively hard particles which abrade the painted surfaces. They are used in polishes to remove the top
20 oxidized layer of the painted surfaces. Therefore, they should be used only occasionally. When these cleaners/polishes of the prior art are used regularly, such abrasive particles cause excessive wear to painted surfaces. The use of cleaners/polishes of the prior
25 art which utilize such abrasives has also been known to damage the vinyl surfaces. A summary of this problem

is discussed in "The Care of Automotive Vinyl Tops," a report of the Vinyl Top Study Task Force, the Chemical Specialties Manufacturers Association, published in Chemical Times & Trends, July 1978, pages 56-57. The abrasives are embedded in the texture of the vinyl, leave an unsightly residue, and mar the vinyl's appearance.

Polymeric solids have been used in cleaning compositions per se. For example, U.S. Pat. No. 4,108,800, issued to Helmut H. Froehlich on August 22, 1978, discloses a cleaning composition wherein polyethylene glycol is added to semi-dry polymeric powdered cleaning compositions to prevent adherence of particles of the cleaning powder to the fabrics being cleaned.

The usefulness of polymeric solids in no-wash-or-rinse water auto cleaner formulations has not been recognized or appreciated in the prior art.

Furthermore, waterless car care products of the prior art such as waxes and cleaners/polishes are instructed to be applied by implements such as cloth, terry towels, or smooth foam pads, and require prior cleaning of the surfaces to remove the soils, lest the soils damage the surfaces.

SUMMARY OF THE INVENTION

According to the present invention there is provided a car cleaner article comprising a dispenser containing a predetermined amount of liquid car cleaner composition and a pad for applying said liquid car cleaner on said car, said pad having resilient fibers and a base, said fibers having a length of from 3 to 15 mm, a diameter of from 10 to 150 microns, and being attached substantially vertically to said base at a density of at least 500 fibers/cm², said pad having a Yield Force of at least 36 Newtons, and said liquid cleaner having:

- I. from 0.1% to 30% by weight of organic polymeric solids selected from
- A. polymeric particles of particle size in the range of from 1 micron to 250 microns;
 - 5 B. polymeric fibers of diameter between 1 micron and 50 microns, and length between 0.1 millimeter to 3 millimeters;
 - C. mixtures of A and B
- 10 said polymeric fibers not exceeding 10% by weight of the composition;
- II. up to 95% of a liquid carrier for said particles; and
- III. an organic suspending agent in sufficient amount to suspend said particles in said liquid carrier.

15 An object of the present invention is to provide a complete car cleaning article which can be used without external source of water. Another object is to provide a substantially nonabrasive liquid car cleaner composition which can be used frequently on car body
20 paint without substantial damage to the paint. Yet another object is to provide an improved vinyl cleaner. Still another object is to provide a resilient fibrous applicator to apply the cleaner composition and scrub soiled car surfaces without letting the soil damage the
25 car surfaces.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a top view of a preferred article comprising a unitary construction of a container in communication with a flocked pad applicator.

30 Fig. 2 is a side view of the article construction of Fig. 1.

Fig. 3 is a planar view of the article viewed from the bottom showing a means of delivering the cleaning composition in the container to the surface
35 to be cleaned.

Fig. 4 shows a puncturing device.

DETAILED DESCRIPTION OF THE INVENTION

The liquid car cleaner composition of this invention comprises organic polymeric solids selected from porous and/or nonporous powdered particles in the particle size range of from 1 micron to 250 microns; and polymeric fibers of diameter between 1 micron and 50 microns, and length between 0.1 millimeter to 3 millimeters. Porous and/or nonporous powdered polymeric particles can be used at a level of 30% by weight of the total composition. A preferred composition contains from 0.5% to 20% of polymeric particles, more preferably from 1% to 10%. But polymeric fibers should be used at a level of no more than 10%. Optimum mixtures of fibers and powders can be formulated in the light of this disclosure. A liquid carrier is required and can be used at a level of up to 95% by weight of the composition. Water and aliphatic hydrocarbon solvents are used as the liquid carrier. Mixtures of water and aliphatic hydrocarbon solvents are preferred. A compatible organic suspending agent in sufficient amount to suspend the particles in the liquid carrier is also required. Both surfactants and thickeners are used as the suspending agent. The surfactants are also used as emulsifier and cleaning aid. Silicone is a preferred optional ingredient and can be used at a level of up to 20% by weight of the composition. Other optional ingredients such as waxes, fluorosurfactants, anticorrosion agents, antistatic agents, sunscreening agents, inorganic mild abrasives, pigments, perfumes, and preservatives can also be used for added benefits.

Polymeric Solids

An essential element of the instant compositions is organic polymeric particulate materials which

are suspended and dispersed throughout the fluid phase. Although the instant invention is not limited to any particular theory or mechanism, it is believed that inclusion of the solid materials in the compositions provides many beneficial effects: (1) promote the uniform spreading and coating of the liquid cleaner on the car surfaces and keep the liquid film uniform (for chemical cleaning) until the cleaner dries off; (2) provide large alternative surface areas to compete with the car surfaces themselves for the soil redeposition (after the soil is lifted up by scrubbing and chemical cleaning actions), when the cleaner is finally dried; (3) act as a soft buffer medium to coat and prevent hard particulate soils from scratching the car surface in this waterless cleaning execution; and (4) spherical-shape particulates provide lubricity by the ball bearing effect. Abrasive solids, when used, provide the polishing action to remove the dead paint layer for surface renewal, but cleaner containing abrasives can only be used occasionally, lest the paint layer is abraded away prematurely. The organic polymeric particulate solids are soft and essentially nonabrasive, therefore the incorporation of these materials in the preferred compositions enables car cleaning without the negative of excessive painted surface wear. Also because the organic polymeric particulates are softer than the common inorganic abrasives, larger size particulates can be used to avoid the deposition of these materials into the depressed areas of the textured vinyl surfaces, without being gritty and surface damaging.

The suitable polymeric particulate materials that can be used are described herein with their overall characteristics. They can be synthetic or naturally-occurring polymeric materials. Synthetic materials which can be utilized include, but are not limited to,

polyethylene, polypropylene, polystyrene, polyester resin, urea-formaldehyde resin, polyvinyl chloride, polyacrylics, polyamide, and copolymers such as ethylene-vinyl acetate copolymer and acrylonitrile-butadiene-styrene terpolymer. Examples of naturally-occurring polymeric materials are cellulosic materials, such as wood powders and short cellulose fibers.

Polymeric particulate materials can be grouped into two general categories, namely, particles (or powders) and short fibers.

The powdery particles can have regular, spherical, or irregular shape. They can be solid or hollow. They can be porous or nonporous. The particle size is substantially in the range of from about 1 micron to about 250 microns.

The nonporous solid particles preferably have spherical shape. They provide both large surface area and lubricity. Some preferred nonporous polymeric particles are polyethylene powders described in "Microthene[®] F Microfine Polyolefin Powders," U.S. Industrial Chemicals Co., Division of National Distillers & Chemical Corp., New York, New York 10016.

The porous particles are made of open cell microporous polymeric materials in which the small void spaces are interconnected. They provide large surface areas for soil deposition. Some preferred porous particles are polyethylene Accurel[®] powder, manufactured by ArmaK Company, Chicago, Illinois 60606 and described in "Versatile Microporous Polymers Developed," Chemical & Engineering News, Vol. 56, Dec. 11, 1978, pages 23-24, and urea-formaldehyde Capture[®] polymer, manufactured by Milliken Chemical, Division of Deering Milliken, Inc., Spartanburg, South Carolina 29304 and described in U.S. Pat. No. 4,194,993, issued to James F. Deal III on March 25, 1980.

Short fiber materials are particulates with elongated forms of diameter between 1 micron to 50 microns, and length between 0.1 millimeter to 3 millimeters. They can be obtained from finely cutting of the fiber filaments. The fibers can be solid or hollow. In the latter case, the fibers have porous property. The preferred short fibers are fibrillated fibers with small fibrils projecting from the surface of the fiber walls. Fibrillated fibers have large surface areas and are believed to have the ability of agglomerating the fine powders and dirt particles. Some preferred fibers are polyethylene Fybrel[®] and Short Stuff[®] fibrillated fibers manufactured by Minifibers, Inc., Weber City, Virginia 24251, and polyethylene and polypropylene Pulpex[®] fibrillated fibers manufactured by Lextar, a Hercules/Solvay Company, Wilmington, Delaware 19899.

The composition of this invention can consist of 0.1% to 30% by weight of porous and/or nonporous polymeric powder particles. A preferred composition of this invention consists of 0.5% to 20% of porous and/or nonporous polymeric particles. A more preferred composition can consist of from 1% to 10% by weight of these particles. Yet another composition of this invention can consist of from 0.1% to 10% by weight of polymeric short fibers. The polymeric solids of this invention can consist of mixtures of powder and fibers, preferably at a ratio of from 20:1 to 1:1 by weight.

Preferred nonporous polymeric powder particles of this invention are: (a) polyethylene of particle size from 5 microns to 150 microns, and used at 0.1% to 30% by weight of the composition; (b) polyethylene of particle size 5 microns to 100 microns, and used at 2% to 15% by weight of the composition; (c) polyethylene particles of particle size 5 microns to 30 microns, and used at 2% to 10% by weight of the composition.

Preferred porous polymeric particles are: (a) urea-formaldehyde polymer 30 microns to 100 microns and used at 0.1% to 30%; (b) polyethylene 30 microns to 150 microns and used at 1.0% to 20%.

5 Preferred fibers are fibrillated polyethylene fibers of: (a) 1 micron to 50 microns in diameter at 0.1 millimeter to 3.0 millimeters in length and used at 0.1% to 10% by weight of the composition; (b) 10
10 microns in diameter at 0.5 millimeter to 1.25 millimeters in length and used at 0.5% to 5% by weight of the composition.

Mixtures of porous and/or nonporous particles and fibrillated fibers are also preferred at level of up to 30% and with amount of fibrillated fibers of not
15 more than 10% by weight of the composition. A more preferred composition consists of a mixture of said polymeric particles and said fibers at a ratio of from 20:1 to 1:1 by weight, and at level of from 0.5% to 20% by weight of the composition.

20

Liquid Carriers

The composition of this invention can contain 2% to 90% by weight of an aliphatic hydrocarbon solvent with boiling points of from 90°C to 300°C or 5% to 95% by weight of water. Liquid carriers comprising mixtures
25 of water and aliphatic hydrocarbons (oil) are preferred. Ratios of 9:1 to 1:9 of water to oil are suitable, and ratios of from 1:1 to 3:1 are preferred. These mixtures are preferably used at 60% to 95%, and more preferably at 70% to 90% by weight of the composition. Preferred
30 amounts of water used in the water-and-oil mixtures are: (a) 30% to 70%; and (b) more preferably 50% to 65% by weight of the total composition.

Preferred aliphatic hydrocarbon solvents are:

(a) Stoddard Solvent, boiling point 160-180°C; (b)
35 Isopar[®] L Solvent (isoparaffinic hydrocarbon solvent produced by Exxon Co., Baltimore, Maryland 21203),

boiling point of 188-207°C; (c) Mineral spirits, boiling point 120-190°C; and (d) Mixture of Stoddard Solvent (160-180°C) and odorless kerosene (190-255°C) at 1:1 to 5:1 weight ratio, all used at 10% to 30%; and more preferably 20% to 30% by weight of the total composition.

Suspending Agents

The suspending agents useful in this invention are suitable surfactants and thickeners and mixtures thereof. These surfactant suspending agents have the properties of dispersing solid particles and liquid droplets. They are used to disperse the polymeric particles throughout the cleaner compositions. Most of the cleaning compositions of this invention contain both oil and water phases. The surfactants also stabilize the emulsion of these two phases. The surfactants are also included to aid in the cleaning of the car surfaces. Substantially any surfactant materials which are compatible with the other components in the composition of this invention can be utilized. These include nonionic, anionic, cationic, amphoteric and zwitterionic surfactants. The composition of this invention can consist of up to 10% by weight of a suspending agent surfactant; preferably between 0.4% and 2%.

The stability of the dispersion and emulsion can also be achieved or further enhanced by addition of a thickener suspending agent to increase the viscosity of the suspending and emulsifying medium.

Thickener suspending agents that can be utilized include, but are not limited to, salts of polyacrylic acid polymer, sodium carboxymethyl cellulose, hydroxyethyl cellulose, acrylic ester polymer, polyacrylamide, polyethylene oxide, natural polysaccharides such as gums, algin, pectins. They are used at effective levels of up to 10%.

Preferred thickeners are salts of polyacrylic acid polymer of high molecular weights. Examples of polyacrylic acid polymers are Carbopol[®] resins which

are described in "Carbopol[®] Water Soluble Resins," Publication No. GC-67, The B. F. Goodrich Co., Cleveland, Ohio 44131. Carbopol[®] resins can be used in the composition of this invention at a level from about 0.05% to about 0.5%, preferably Carbopol[®] 934 used at 0.1% to 0.2% by weight of the total composition. Sodium hydroxide and other inorganic and organic bases are utilized in the compositions of this invention at effective levels to neutralize the Carbopol[®] thickeners, as described in the publication mentioned above.

A preferred thickener suspending agent which is utilized in nonaqueous compositions is glyceryl tris-12-hydroxystearate manufactured under the name of Thixcin R[®] by NL Industries, used preferably in the range of from 0.2% to 2% by weight of the total composition.

Optional Ingredients

Compatible optional ingredients can be used in the composition of this invention for added benefits. Silicone is a preferred optional component. Silicone materials provide or enhance the gloss/shine appearance of car surfaces, improve the ease of application and removal of the cleaner, and make the car surfaces water repellent for added protection. Silicone materials which can be used include, but are not limited to, dimethyl silicones, aminosilicones, silicone resins, and mixtures thereof. Preferred silicones are the dimethyl silicones and aminosilicones. Examples of dimethyl silicones are the Dow Corning[®] 200 Fluids of various viscosities, manufactured by Dow Corning Corp., Midland, Michigan 48640. Examples of aminosilicones are the Dow Corning[®] 531 and 536 Fluids. These Dow Corning[®] Fluids will be referred to hereinafter by the abbreviated name "DC". Silicone materials can be used in the composition of this invention at a level of up to 20%. Preferred silicone materials and levels are:

(a) DC-200, viscosity 50-10,000 centistokes, used at 1%

to 10%; (b) DC-200, viscosity 100-1000 centistokes, used at 2% to 6%; and (c) mixture of DC-531 and DC-536 at 3:1 to 6:1 weight ratio, and at 1% to 10% by weight of the total composition.

5 Other optional ingredients that can be used in the composition of this invention include, but are not limited to, waxes for surface protection, fluorosurfactants for spreadability and leveling, other organic solvents for greasy soil cleaning, anticorrosion agents,
10 antistatic agents, pigments, perfumes, preservatives.

Mild inorganic abrasives such as calcium carbonate powder can also be used when polishing action is desired so long as they do not leave unsightly residue on textured vinyl surfaces.

15 Dispenser and Applicator

In this dry cleaning execution it is essential that the application implement has a construction such that it: (1) provides effective spreading and scrubbing, resulting in good cleaning and uniform end result appearance on painted surfaces; (2) prevents the gritty soil
20 particles from incurring scratches to the painted surfaces; and (3) can reach to dislodge the embedded soil in the depressed areas of the textured vinyl surfaces.

It was discovered that a bristle-fibered pad
25 with the defined fiber construction (as described herein) can be used to apply the active composition to clean soiled car painted surfaces virtually without damaging those surfaces. Although the instant invention is not limited to any particular theory or mechanism, it is
30 believed that the bristle-fibered application pad provides the desired properties for surface-safe cleaning because: (1) It has enough void volume to hold the gritty soil particles and to keep them away from the car surfaces, thus preventing them from scratching the car
35 surfaces; (2) It has vertical fibers that stay essentially unbent under normal hand scrubbing pressure to

keep the gritty soil particles in the void spaces and away from the car surfaces (long and/or thin fibers bend under this pressure and push some gritty particles onto the surface); (3) It has straight vertical fibers which
5 can reach depressed areas of the textured vinyl surface; and (4) It has high surface fiber density (number of fibers per unit area) to provide effective scrubbing and cleaning for good end result appearance.

The applicator/scrubbing pad is constructed
10 essentially of bristled fibers secured vertically to a base. Flocking is a preferred method of fiber attachment. In this preferred method, the fibers are attached to the base by electrostatic flocking for good vertical fiber alignment, using a flocking adhesive such as an
15 acrylic adhesive made from Rhoplex[®] resin manufactured by Rohm and Haas Co., Philadelphia, Pennsylvania 19105. Tufting is also a preferred method of fiber attachment: pile fabric which consists of fibers vertically tufted into a woven yarn substrate. The fabric is then adhes-
20 ively laminated to the base. The fibers are made of resilient polymeric materials, preferably nylon, polypropylene, acrylic, modacrylic, polyester.

Following are the requirements of fiber composition and pad construction for a good performing
25 applicator/scrubbing pad:

1. Fiber density of at least 500 fibers/cm² to provide effective scrubbing and cleaning.
2. Said fibers have a minimum fiber length of 3 mm so that they can reach to scrub and
30 clean the depressed areas of the textured vinyl surfaces.
3. Said applicator/scrubbing pad must have a large enough surface area for fast cleaner application and scrubbing of the total car exterior surfaces. The pad surface area
35 should be at least 60 cm².

4. The fibers must be aligned substantially vertically to the base, and the fibers must remain essentially unbent under normal hand scrubbing pressure.

5 The last requirement above can be defined by the "Yield Force" which is the minimum force needed to bend the fibers of the pad. The Yield Force of the pad must be greater than the normal hand scrubbing force of 22-36 Newtons (5-8 lbs.). The Yield Force of a pad
10 is a collective property affected by many factors, which include fiber material, fiber length, fiber diameter, fiber density, fiber orientation (relative to base), nature of the base, and effective pad surface area. The Yield Force of a pad can be measured directly with an
15 Instron tester (see below), or calculated from the "Yield Pressure" and the pad surface area by the relation:

Yield Force = Yield Pressure x pad surface area.

Yield Pressure is the minimum force exerted
20 vertically upon a unit area of the pad to bend the fibers.

Yield Pressure is determined by the same procedure of the Compression Test as described in the standard method ASTM D-695 by using an Instron tester, Model TM, manufactured by the Instron Corp., Canton,
25 Massachusetts 02021. A fibrous pad cut to a predetermined effective surface area A is placed on the compression cell of the Instron tester. Test specimens of square or circular form with effective surface area of between 58 cm² and 180 cm² are recommended. Testing speed of 0.51 cm/min.
30 (0.2 in./min.) is recommended. The force F required to bend the fibers is read from the load indicator recording chart. The Yield Pressure is the ratio F/A.

Examples of fibrous materials used in the construction of the applicator/scrubbing pad of this
35 invention are listed in Table 1. The Yield Pressures of these materials are listed in Table 2.

TABLE 1

Ex.	Pad Surface	Fiber Length (mm)	Fiber Diameter (μm)	Fiber Density (fib/cm ²)	Fiber Mat'l	Others
5	1 Padco [®]	4.6	47	1880	Nylon	a,c,g
	2 Padco [®]	4.6	47	1880	Nylon	a,d,g
	3 Padco [®]	4.6	47	1880	Nylon	a,e,g
	4 IF-455	5.6	43	3570	PP ⁱ	b,f,h
	5 IF-456	5.1	43	3570	PP ⁱ	b,f,h
10	6 IF-457	4.6	43	3570	PP ⁱ	b,f,h
	7 IF-458	4.1	43	3570	PP ⁱ	b,f,h
	8 Scrubber [®]	5.6	49	8120	PP ⁱ	b,f,h
	9 IF-498	5.8	44	2970	Nylon	b,f,h
	10 IF-507	6.9	44	2970	Nylon	b,f,h

15 Method of attachment of fibers to base:

(a) flocked

(b) tufted

Base construction:

- (c) polyurethane foam, 1.6 mm thick
- (d) polyurethane foam, 4.8 mm thick
- (e) polyurethane foam, 7.9 mm thick
- (f) woven yarn

Manufacturers:

- (g) Padco, Inc., Minneapolis, Minnesota 55414;
- (h) Collins & Ackman Corp., Roxboro, North Carolina 27573.

Fiber Material:

(i) PP = polypropylene

TABLE 2

Ex.	Pad Surface (Described in Table 1)	Yield Pressure	
		(psi)	($\times 10^3 \text{ N/m}^2$)
5	1 Padco ^(R)	1.05	7.2
	2 Padco ^(R)	0.80	5.5
	3 Padco ^(R)	0.65	4.5
	4 IF-455	1.75	12.1
	5 IF-456	2.35	16.2
10	6 IF-457	3.85	26.5
	7 IF-458	3.85	26.5
	8 Scrubber ^(R)	10.00	69.0
	9 IF-498	2.00	13.8
	10 IF-507	1.70	11.7

15 To calculate the Yield Force of an applicator/
scrubbing pad, one first determines the Yield Pressure
of the fibrous material and the desired effective surface area
the pad, then takes the product of the two values. An
acceptable applicator/scrubbing pad of this invention
20 must have a Yield Force greater than the normal hand
scrubbing force of 36 Newtons (8 lbs.). Example: An
applicator/scrubbing pad, with an effective surface area of 116
 cm^2 (18 in.^2) and constructed with Padco flocked material
with 4.8 mm thick polyurethane foam base (Example 2 of
25 Table 1) has a Yield Force of 64 Newtons (14.4 lbs.)
which is greater than 36 Newtons, therefore satisfies
the requirement number 4 above.

Preferably fibers have length of from 3 mm to
15 mm and diameter of from 10 microns to 150 microns.
30 Fiber density is at least 500 fibers/ cm^2 ; more prefer-
ably at least 1500 fibers/ cm^2 . Examples of flocked
and tufted materials that can be used for the applicator/
scrubbing pad of this invention are listed in Table 1.

The base of the applicator/scrubbing pad can be a foam pad or a semi-rigid but flexible plastic film. The preferred base is a close-cell foam pad with fine pores, preferably more than 20 pores per linear centimeter. A preferred foam pad is made of close-cell polyurethane foam with 28-32 pores per linear centimeter. Preferred foam thickness is from 1 mm to 10 mm.

Preferably the fibers cover the total application surface (i.e. the effective surface area) of the pad. The pad has a minimum effective surface area of 60 cm^2 , preferably from 100 cm^2 to 200 cm^2 . A more preferred pad has a dimensions of about 8 cm x 20 cm. Preferably it has one long end tapered into a point to enable the pad to clean tight spots, as depicted in Figs. 1 and 3.

The dispenser can be made of any materials which are compatible with the cleaner composition, such as metal or plastic materials, preferably polyethylene and polypropylene. The dispenser preferably has a palm-fitting shape with resilient side walls. The dispenser has opening means for cleaner loading and dispensing. In a preferred construction the dispenser has a dispensing valve such as a diaphragm valve described in U.S. Pat. No. 4,226,342, issued to Robert H. Laauwe on October 7, 1980, or a duckbill valve available from Vernay Laboratories, Inc., Yellow Springs, Ohio 45387.

For convenience, it is preferable that the dispenser and the pad are of a unitary construction, in which a palm-fitting container holding a predetermined amount of liquid cleaner composition is positioned on top of the applicator/scrubbing base with a means to dispense the liquid cleaner to the car surfaces. The dispensing means can be an aperture opening through the applicator pad or at the tip of the pad. The aperture can be sealed initially with a thin plastic film which is punctured to discharge the cleaner. For the through-the-pad dispensing method, the dispensing aperture can

be adapted with a diaphragm valve. For the through-the-tip dispensing method, the dispensing aperture can be adapted with a duckbill valve. Preferably the container has a capacity and contains of from 150 5 cm³ to 300 cm³ of the liquid cleaner.

DESCRIPTION OF THE DRAWINGS

Figs. 1 and 2 show, respectively, top and side views of a preferred dispenser/applicator article which is used to apply the liquid cleaner. Fig. 3 is a bottom view of the article. Fig. 4 shows a puncturing device. This dispenser/applicator article comprises: an applicator/scrubbing pad 1 and a container 2 which contains the liquid cleaner 20.

The applicator/scrubbing pad 1 has bristle 15 flocked fibers 3 secured to a base 4 with an acrylic flocking adhesive 4a. Pad 1 has a slit opening 5 as shown in Fig. 3. The pad has dimensions of about 8 cm x 20 cm. It has one long end 21 tapered into a point to enable the pad to clean tight spots, as depicted in 20 Fig. 3. The pad construction consists of nylon fibers 3 of 4.6 cm length, 47 microns diameter (18 denier) flocked onto a close-cell foam pad 4 to a density of about 1900 fibers/cm² (172 g/m²). The foam pad 4 is made of close-cell polyurethane foam with 28-32 pores 25 per linear centimeter. The foam pad 4 has a thickness of 4.8 cm. The foam is attached to a semi-rigid plastic base 6 by means of a suitable adhesive 7, such as a hot melt adhesive. The container 2 has a palmfitting shape with resilient side walls 18. The container 2 is made 30 by blow molding polyethylene. The container has a capacity of 230 cm³. The container 2 is positioned on top of the base 6 by close-fitting annular projections 8 into the openings 10 in base 6. The container 2 is secured to the base 6 by using a suitable adhesive 12.

such as a hot melt adhesive. The container 2 has an aperture means 9 through which the cleaner 20 will be dispensed. This aperture 9 is aligned with the opening 11 of the base 6 and the opening 5 of the pad 1. The aperture 9 is initially sealed off by a thin plastic film 13, such as a pressure sensitive tape. The reservoir 2 also has an opening 14, with circumferential groove (not shown) and a screw cap means 16, via which the cleaner 20 is loaded or refilled.

10 To discharge the cleaner 20, the sealing film 13 is first punctured via opening 5 using a sharp puncturing device as shown in Fig. 4. Then the cleaner can be discharged from the container to the surface to be cleaned via the aperture 9 by hand pressure to the side walls 18.

It will be understood that other embodiments of the dispenser/applicator article come within the scope of this disclosure, e.g., the bristled pad can take the form of a mitten made of flocked material and the liquid cleaner can be in a separate plastic bottle dispenser.

End Result Appearance Performance Test

This is a test method to evaluate the end result appearance performance of the cleaner compositions on painted and textured vinyl surfaces. A composition is considered acceptable if after use (as described below), it leaves a uniform appearance on painted surfaces, i.e., substantially free of streaks, and does not leave any appreciable amount of unsightly residue embedded in the texture of the vinyl surface.

Test Procedure

Test painted surfaces are black acrylic enamel painted plates of dimension 30.5 cm x 30.5 cm. Test vinyl surfaces are textured vinyl sheets of dimension 30.5 cm x 30.5 cm. This car top vinyl material with Milano grain pattern, color M398 (Midnite Blue), is manufactured by Weymouth Art Leather Co., South Braintree, Massachusetts 02184.

The cleaner compositions are applied to the test surfaces with fiber-flocked foam pads of 5.1 cm x 7.6 cm pad surface dimensions. These pads are comprised of a close-cell polyurethane foam base of 9.5 millimeters thickness and flocked with 18 denier nylon fibers of 4.6 mm fiber length, and flock density of 172 g/m². The pad material is manufactured by Padco, Inc., Minneapolis, Minnesota 55414.

All cleaning tests are performed in a laboratory with controlled temperature and humidity conditions, namely, 27°C and 15% relative humidity.

Two milliliters of a cleaner composition is dispensed to the test surface and spread with a fiber-flocked foam pad to cover the surface with a circular rubbing motion. The cleaner is let dry to a powdery haze, then the haze is wiped off with a terry cloth, and the surface appearance evaluated.

The following examples are given for purposes of illustration only and are not to be interpreted as necessarily limiting the invention. All percentages are by weight unless otherwise indicated.

EXAMPLE I

	<u>Raw Materials & Source</u>	<u>Wt. %</u>	<u>Chemical Description</u>
5	1. Stoddard Solvent (Fisher)	26.0	Petroleum distillates (b.p. 153-210°C)
	2. DC-200 [®] Silicones (350 cts) (Dow Corning)	4.0	Polydimethylsiloxane
10	3. Calamide [®] C Surfactant (Pilot Chemical)	1.0	Coconut diethanol- amide
	4. Carbopol [®] 934 Thickener (2% solution) (B.F. Goodrich)	10.0*	Polyacrylic acid polymer
15	5. Deionized Water	50.95*	
	6. Sodium Hydroxide (10% solution)	1.05*	
20	7. Short Stuff [®] 13040F Fibers (Minifibers)	1.0	Polyethylene fibril- lated fibers (0.8- 1.05mm fiber length)
	8. Microthene [®] FA-520 Powder (USI Chemicals)	4.0	Polyethylene powder (20 micron particle size)
25	9. Capture [®] Polymer (Milliken Chemicals)	2.0	Urea-Formaldehyde porous powder (40- 110 micron particle size)

30 Total 100.00%

*Total water is 60.695%.

Preparation Directions for Example I

- Step I: Add 2 and 3 to 1 with stirring.
- Step II: Separately prepare solution 4 and solution 6.
- Step III: Add 4 to 5 with continuous stirring.
- 5 Step IV: Add 6 to the mixture of Step III with good stirring until the mixture thickens uniformly.
- Step V: Add the mixture of Step I to the mixture of Step IV with continuous stirring to
- 10 form a thick, smooth, creamy emulsion.
- Step VI: Add 7, 8 and 9, in that order, to the mixture of Step V with continuous stirring until all are well dispersed.

The composition of Example I contains a total

15 of about 7% polymeric solids, 87% liquid carrier, 1.3% suspending agents and 4% silicone. End Result Appearance Tests showed that the composition of Example I is acceptable for painted and vinyl surfaces.

The composition of Example I requires no

20 prewashing or rinsing of car surface before use. However, one may wish to remove heavy soil such as caked mud prior to using the product. The product is good for cleaning most exterior car surfaces. For best results, user should avoid direct sunlight and allow

25 car to cool before use.

Usage Instructions

1. Shake the cleaner to assure uniformity.
2. Apply on car surfaces, preferably with the container/applicator kit as shown in the drawings.
- 30 3. Start at the top of car and work down. Spread product to cover surface with a uniform film. Rub with circular motion.

4. Let product dry to powdery haze, loosened dirt and grime will be trapped as the product dries to a powdery haze.
5. Wipe off haze with clean cloths and turn cloths frequently.

Other Examples:

In general, the compositions of the following Examples were made by following the procedure of Example I, namely, by: (1) mixing the silicone and the surfactant into the organic solvent (oil) phase, (2) mixing the Carbopol thickener and neutralizers into the water phase, (3) mixing the oil phase into the water phase, and (4) adding the polymeric particulate solids to the liquid emulsion with continuous stirring until they are uniformly dispersed. Any variations to this procedure are noted under the appropriate Examples. The preferred order of addition of the particulate solids is fibers first, then nonporous particles, and finally porous particles. High shear mixing for a short period of time after all ingredients have been added is preferred in order to break up any clumping of the solid materials, and to achieve thorough mixing.

	<u>Ingredients</u>	<u>Ex. II</u>	<u>Ex. III*</u>
	Microthene [®] FA-520 Polyethylene Powder (USI Chemicals)	6.0	-
5	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	2.0	-
10	Short Stuff [®] 13040 Poly- ethylene Fibrillated Fibers (Minifibers)	1.0	-
	Snowflow [®] Diatomaceous Silica (Johns Manville)	-	9.0
	Stoddard Solvent Petroleum Distillates	26.0	26.0
15	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0
	Oleic Acid	1.0	1.0
20	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	5.0	5.0
	Triethanolamine (2% solution)	5.0	5.0
	Morpholine	0.6	0.6
	Deionized Water	<u>49.4</u>	<u>49.4</u>
	Totals	100.0	100.0
25	Residue on Vinyl	No	Heavy

*Outside scope of the present invention.

Example II: Procedure of Example I, except that the fibrillated fibers are added to the water phase.

30 Example III: Procedure of Example I, with both neutralizers, namely, triethanolamine and morpholine, are added to the water phase, and oleic acid is added to the oil phase.

	<u>Ingredients</u>	<u>Ex. IV*</u>	<u>Ex. V</u>
	Microthene [®] FA-520 Polyethylene Powder (USI Chemicals)	-	4.0
5	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	-	2.0
10	Short Stuff [®] 13040 Poly- ethylene Fibrillated Fibers (Minifibers)	-	1.0
	Snowflow [®] Diatomaceous Silica (Johns Manville)	9.0	-
15	Gelwhite [®] GP Montmorillonite Clay (8% dispersion) (Georgia Kaolin)	12.5	-
	Stoddard Solvent Petroleum Distillates	26.0	26.0
	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0
20	Oleic Acid	1.0	-
	Dodecylamine	-	0.2
	Dimethyldodecylamine	-	0.2
25	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	-	10.0
	Sodium Hydroxide (10% solution)	-	0.8
	Morpholine	0.6	-
	Deionized Water	<u>46.9</u>	<u>51.8</u>
	Totals	100.0	100.0
30	Residue on Vinyl	Heavy	No

*Outside scope of the present invention.

0100195

Example IV: Add clay, diatomaceous silica and morpholine to the water phase, and oleic acid to the oil phase.

Example V: Procedure of Example I, with the amines added to the oil phase.

	<u>Ingredients</u>	<u>Ex. VI</u>	<u>Ex. VII</u>
	Microthene [®] FA-520 Polyethylene Powder (USI Chemicals)	29.0	10.0
10	Stoddard Solvent Petroleum Distillates	26.0	26.0
	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0
15	Calamide [®] C Cocodiethanolamide (Pilot Chemical)	1.0	1.0
	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0
20	Sodium Hydroxide (10% solution)	1.05	1.05
	Deionized Water	<u>28.95.</u>	<u>47.95</u>
	Totals	100.00	100.00
	Residue on Vinyl	Slight	No

25 Example VI: Add half of the solids to the water phase, the other half to the oil phase, then add oil phase to water phase.

<u>Ingredients</u>		<u>Ex. VIII</u>	<u>Ex. IX</u>
	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	29.0	18.0
5	Stoddard Solvent Petroleum Distillates	26.0	26.0
	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0
10	Calamide [®] C Cocodiethanolamide (Pilot Chemical)	1.0	0.5
	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0
15	Sodium Hydroxide (10% solution)	1.05	1.05
	Deionized Water	<u>28.95</u>	<u>40.45</u>
	Totals	100.00	100.00
	Residue on Vinyl	No	No

20 Example VIII: Procedure of Example VI.

0100195

	<u>Ingredients</u>	<u>Ex. X</u>	<u>Ex. XI</u>
	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	8.0	-
5	Accurel [®] Polyethylene Porous Powder (Armak)	-	7.0
	Stoddard Solvent Petroleum Distillates	26.0	26.0
10	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0
	Calamide [®] C Cocodiethanolamide (Pilot Chemical)	0.5	0.5
15	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0
	Sodium Hydroxide (10% solution)	1.05	1.05
	Deionized Water	<u>50.45</u>	<u>51.45</u>
20	Totals	100.00	100.00
	Residue on Vinyl	No	No

Example XI: Procedure of Example I, except the solid powder is added to the oil phase.

Ingredients		Ex. XII	Ex. XIII	Ex. XIV
	Short Stuff [®] 13040 Polyethylene Fibrillated Fibers (Minifibers)	10.0	7.0	3.0
5	Stoddard Solvent Petroleum Distillates	26.0	26.0	26.0
	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0	4.0
10	Calamide [®] C Cocodiethanolamide (Pilot Chemical)	1.0	0.5	0.5
	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0	10.0
15	Sodium Hydroxide (10% solution)	1.05	1.05	1.05
	Deionized Water	<u>47.95</u>	<u>51.45</u>	<u>55.45</u>
	Totals	100.00	100.00	100.00
	Residue on Vinyl	No	No	No

20 Examples XII, XIII, and XIV: Procedure of Example XI.

0100195

	<u>Ingredients</u>	<u>Ex. XV</u>	<u>Ex. XVI</u>	<u>Ex. XVII</u>
	Microthene [®] FA-520 Polyethylene Powder (USI Chemicals)	5.0	8.0	-
5	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	4.0	-	8.0
10	Short Stuff [®] 13040 Poly- ethylene Fibrillated Fibers (Minifibers)	-	1.0	1.0
	Stoddard Solvent Petroleum Distillates	26.0	26.0	26.0
	DC-200 [®] , 350 cts (Dow-Corning)	4.0	4.0	4.0
15	Dodecylamine	0.2	0.2	0.2
	Dimethyldodecylamine	0.2	0.2	0.2
	Carbopol [®] 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0	10.0
20	Sodium Hydroxide (10% solution)	0.52	0.52	0.52
	Deionized Water	<u>50.08</u>	<u>50.08</u>	<u>50.08</u>
	Totals	100.00	100.00	100.00
	Residue on Vinyl	No	No	No

25 Example XVI: Procedure of Example I, except the fibers
are added to the water phase.

0100195

	<u>Ingredients</u>	<u>Ex.XVIII</u>	<u>Ex.XIX</u>	<u>Ex. XX</u>
	Microthene ^(R) FA-520 Polyethylene Powder (USI Chemicals)	5.0	4.0	8.5
5	Capture ^(R) Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	3.0	2.0	2.0
	Accurel ^(R) Polyethylene Porous Powder (Armak)	-	2.0	-
10	Short Stuff ^(R) 13040 Poly- ethylene Fibrillated Fibers (Minifibers)	1.0	1.0	0.5
	Stoddard Solvent Petroleum Distillates	26.0	26.0	19.5
15	Kerosene (Deodorized)	-	-	6.5
	DC-200 ^(R) , 350 cts (Dow-Corning)	4.0	4.0	4.0
20	Calamide ^(R) C Cocodiethanolamide (Pilot Chemical)	-	-	1.0
	Dodecylamine	0.2	0.2	-
	Dimethyldodecylamine	0.2	0.2	-
25	Carbopol ^(R) 934 Polyacrylic Acid Resin (2% solution) (B.F. Goodrich)	10.0	10.0	5.0
	Sodium Hydroxide (10% solution)	0.52	0.52	0.53
	Deionized Water	<u>50.08</u>	<u>50.08</u>	<u>52.47.</u>
	Totals	100.00	100.00	100.00
30	Residue on Vinyl	No	No	No

Example XIX: All particles are added to the water phase, fibers to the oil phase, then add oil phase to water phase.

	<u>Ingredients</u>	<u>Ex. XXI</u>	<u>Ex. XXII</u>
	Microthene [®] FA-520 Polyethylene Powder (USI Chemicals)	4.0	4.0
5	Capture [®] Polymer Urea- Formaldehyde Porous Powder (Milliken Chemicals)	2.0	-
	Accurel [®] Polyethylene Porous Powder (Armak)	-	5.5
10	Short Stuff [®] 13038F Poly- ethylene Fibrillated Fibers (Minifibers)	1.0	0.5
	Stoddard Solvent Petroleum Distillates	-	89.0
15	Thixcin R [®] Glyceryl tris-12-hydroxystearate suspending agent (NL Industries)	-	1.0
20	Carbopol [®] 940 Polyacrylic Acid Resin (1% aqueous solution) (B.F. Goodrich)	5.0	-
	Sodium Hydroxide (10% solution)	0.26	-
	Deionized Water	<u>87.74</u>	<u>-</u>
25	Totals	100.00	100.00
	Residue on Vinyl	No	Very slight

Example XXII: (1) Warm the Stoddard Solvent to 50°C
in a water bath; (2) sprinkle Thixcin R[®] into the
Stoddard Solvent (still in the water bath) with
30 vigorous stirring using a cutting blade paddle;
(3) the mixture is subjected to high sheer mixing;
(4) add the solids with continuous stirring; (5) the
final composition (at 50°C) is subjected to high sheer
mixing; and (6) stir the mixture with a cutting blade
35 paddle until cooled down to room temperature.

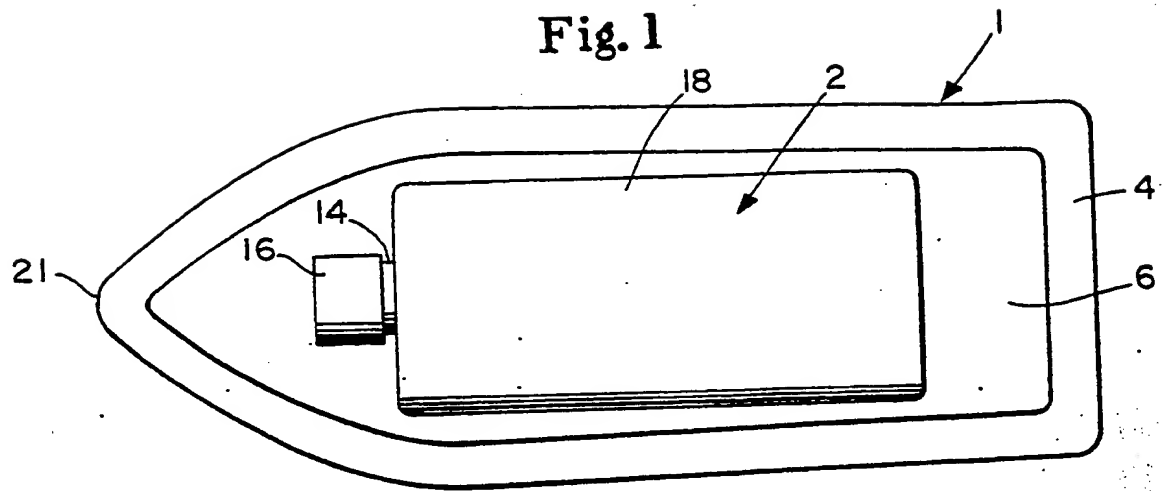
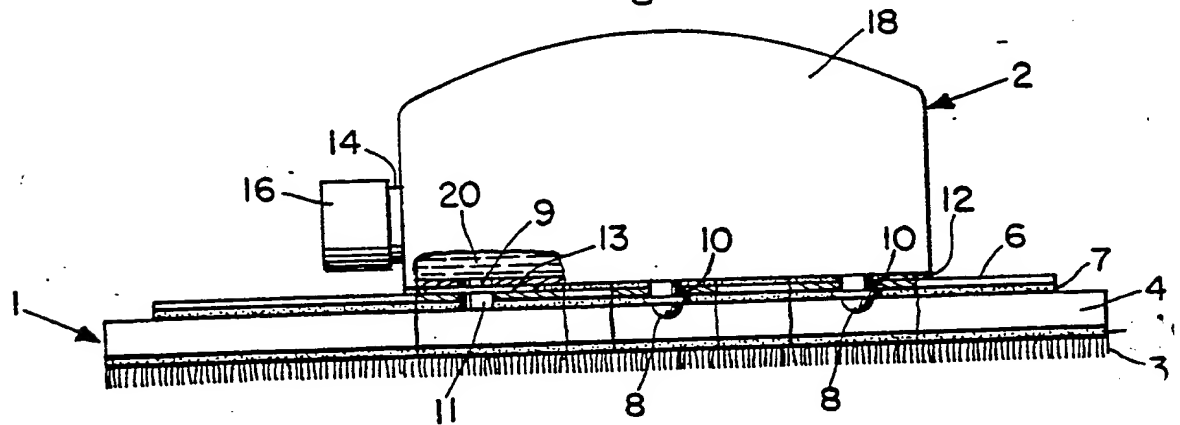
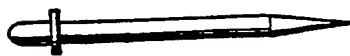
Compositions of Examples III and IV which contain diatomaceous silica abrasives, and clay and diatomaceous silica abrasives, respectively, left heavy residues on vinyl surfaces according to the End Result 5 Appearance Performance Test, and fall outside the scope of the present invention. Compositions of all other Examples contain organic polymeric particulates, left no residue or only very small amount of residues, fall within the scope of this invention.

CLAIMS:

1. A car cleaner article comprising a dispenser containing a predetermined amount of liquid car cleaner composition and a pad for applying said liquid car cleaner on said car, said pad having resilient fibers and a base, said fibers having a length of from 3 to 15 mm, a diameter of from 10 to 150 microns, and being attached substantially vertically to said base at a density of at least 500 fibers/cm², said pad having a Yield Force of at least 36 Newtons, and said liquid cleaner having:
 - I. from 0.1% to 30% by weight of organic polymeric solids selected from
 - A. polymeric particles of particle size in the range of from 1 micron to 250 microns;
 - B. polymeric fibers of diameter between 1 micron and 50 microns, and length between 0.1 millimeter to 3 millimeters;
 - C. mixtures of A and B
 - II. up to 95% of a liquid carrier for said particles; and
 - III. an organic suspending agent in sufficient amount to suspend said particles in said liquid carrier.
2. An article according to Claim 1 wherein said pad comprises fibers having a length of from 4 mm to 8 mm, a diameter of from 30 microns to 60 microns, and a fiber density of at least 1200 fibers/cm².
3. An article according to either one of Claims 1 and 2 wherein said dispenser and pad are of a unitary construction, being formed with means to dispense said liquid cleaner to car surfaces.

4. An article according to any one of Claims 1-3 wherein said container contains from 150 cm³ to 300 cm³ of said liquid cleaner.
5. An article according to any one of Claims 1-4 wherein said pad has a surface area of from 100 cm² to 200 cm².

1/1

Fig. 1**Fig. 2****Fig. 4****Fig. 3**